05/16/2006 14:35 2128080844 NORRIS MCLAUGHLIN PAGE 02/09

AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the

application:

Listing of Claims:

1. (Currently Amended) Pharmaceutical preparation with retarding

active ingredient release, comprising a mixture of powdery teicoplanin and at least one

powdery, water soluble salt form of at least one of gentamicin, clindamycin, kanamycin,

amikacin, tobramycin, vancomycin, moxifloxacin and ciprofloxacin and an inorganic

and/or organic adjuvant.

2. (Currently Amended) Pharmaceutical preparation with retarding

active ingredient release pursuant to claim 1, which contains calcium carbonate, calcium

sulfate dihydrate, tricalcium phosphate and/or hydroxylapatite as the inorganic adjuvant.

3. (Currently Amended) Pharmaceutical preparation with retarding

active ingredient release pursuant to claim 1, which contains polyesters of at least one of

lactic acid, glycolic acid, 5-hydroxy valeric acid, 6-hydroxy caproic acid and co-polymers

thereof as organic adjuvants.

4. (Currently Amended) Pharmaceutical preparation with retarding

active ingredient release pursuant to claim 1, which is in the form of one of tablets,

molded bodies, fibers and granules.

USSN 10/600,557

Amendment under 37 CFR § 1.111 filed May 16, 2006

- 5. (Currently Amended) Pharmaceutical preparation with retarding active ingredient release pursuant to claim 1, comprising a combination of polymerizable methacrylic acid esters and mixtures consisting of powdery teicoplanin and at least one powdery, water soluble salt form of at least one of gentamicin, clindamycin, kanamycin, amikacin, tobramycin, vancomycin, moxifloxacin and ciprofloxacin formed and polymerized into a molded body.
- 6. (Currently Amended) Pharmaceutical preparation with retarding active ingredient release pursuant to claim 1, wherein the mixture is part of a resorbable and/or of non-resorbable coating, which has been applied to non-metallic and metallic implants.
- 7. (Currently Amended) Pharmaceutical preparation with retarding active ingredient release pursuant to claim 1, wherein before being cured inorganic calcium phosphate bone cements and plaster mixtures are admixed to mixtures consisting of powdery teicoplanin and at least one powdery, water soluble salt form of at least one of gentamicin, clindamycin, kanamycin, amikacin, tobramycin, vancomycin, moxifloxacin and/or ciprofloxacin.
- 8. (Currently Amended) A permanent or temporary implant comprising a pharmaceutical preparation with retarding active ingredient release pursuant to claim 1 in the form of one of tablets, molded bodies, fibers and granules.

USSN 10/600,557 3 Amendment under 37 CFR § 1.111 filed May 16, 2006 9. (New) A method of treating an antibacterial infection in a patient in need thereof comprising administering to said patient a pharmaceutical preparation pursuant to claim 1.

USSN 10/600,557 4
Amendment under 37 CFR § 1.111 filed May 16, 2006